

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

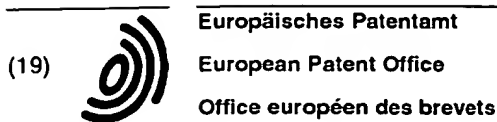
Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

**As rescanning documents *will not* correct images,
please do not report the images to the
Image Problem Mailbox.**



(11) **EP 0 908 558 A1**

(12) **EUROPEAN PATENT APPLICATION**

(43) Date of publication:
14.04.1999 Bulletin 1999/15

(51) Int Cl.⁶: **D21H 11/10, D21H 23/72**

(21) Application number: **98660095.5**

(22) Date of filing: **16.09.1998**

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

- Niskanen, Kaarlo Johannes
06150 Porvoo (FI)
- Nygard, Stina
08500 Lohja as. (FI)
- Pitkänen, Malja
40250 Jyväskylä (FI)

(30) Priority: **16.09.1997 FI 973706**

(71) Applicant: **METSÄ-SERLA OY**
02100 Espoo (FI)

(74) Representative: **Sundman, Christoffer et al**
Seppo Laine Oy,
Itämerenkatu 3 B
00180 Helsinki (FI)

(72) Inventors:
• Leskelä, Markku
08500 Lohja as. (FI)

(54) **Process for preparing base paper for coated fine paper**

(57) The present invention concerns a method for producing a base paper for fine papers, the base paper being produced from a mixture of a mechanical pulp and a chemical pulp. The chemical pulp used comprises a chemical softwood pulp having a elastic modulus close to that of the mechanical pulp and a great bonding strength. It is preferred to use a chemical softwood pulp

which produces a sheet having a elastic modulus of less than 6000 N/mm² when the bonding strength is 400 J/m². A pulp of this kind has a good ScottBond strength at the same light scattering. The base paper produced by the method can therefore be used in double-coated fine papers which require a large bonding strength of the base paper.

EP 0 908 558 A1

Description

[0001] The present invention concerns a process according to the preamble of claim 1 for preparing a base paper that can be used as base paper in particular for manufacturing coated fine papers or printing paperboards. A paper or paperboard of this kind comprises bleached chemical pulp.

[0002] The present invention also concerns a base paper for fine papers according to the preamble of claim 13 and a process according to the preamble of claim 14 for manufacturing fine paper.

[0003] The special problem of coated, in particular double-coated, fine papers is that the paper web tends to split in the dryer of the printing machine when water from the printing colour and similar solvents are removed by drying. The problem is caused by the fact that double-coating forms on the surface of the paper a very dense coating layer which cannot be penetrated by steam vapourizing from the base paper. The steam primarily stems from the normal 4 to 5 % moisture content of paper and the bubbles formed from the moisture break the paper, if the strength properties of the base paper are not sufficient for resisting this steam pressure.

[0004] The afore-described problem is called blistering and the required internal bond strength (z-directional strength) of the paper is measured by the ScottBond value.

[0005] Traditionally, a reduction of the blistering of the base paper of fine papers has been aimed at by increasing the beating of the chemical pulp, in order to obtain more bonds between the fibers. This solution comprises the disadvantage that an increase of the beating does not enhance the bonding strength expressed by the ratio of strength-to-bonding surface area. Increased beating causes a number of problems. First, when the beating is increased, dewatering of paper is impaired. Therefore, the water content of the paper is disadvantageously high when the paper after web forming is transferred to the wet press section of the paper machine and then onwards to the drying section. As a result, it becomes more likely that the paper will adhere to the rollers of the wet press and drying sections, and the risk of web breaks increases. Further, the strength of the web is small at higher water contents and this already increases the risk of web breaks.

[0006] Secondly, also the properties of dry paper change in an undesirable way if the pulp is subjected to extensive beating. When the beating is increased the density of the paper grows and as a result the stiffness of the paper decreases. This causes runability problems in the paper machine due to wavy edges. When paper density grows, the fibers of the chemical pulp are more and more tightly bonded so that the elastic modulus increases. Then the paper becomes brittle and its toughness is not sufficient to meet the strain caused by the paper and printing machines.

[0007] It should be mentioned that the insufficient internal bond strength of paper causes problems also during sheet offset printing although no separate dryer is used in that printing technique. In sheet offset printing the problem is formed because the printing colours are sticky. When the paper is released from the printing nip, the surface of the paper and the wet printing colour are stuck together. If the internal bond strength of the paper is not large enough in comparison to the internal cohesion forces of the printing colour, the surface of the paper will accompany the printing colour and the paper will split in the middle of the sheet. Increased beating of the chemical pulp has been used in attempts to solve this problem also.

[0008] It is an object of the present invention to eliminate the problems of the prior art and to provide an entirely novel method for producing a paper web which can be used as a base web for coated fine papers. In particular it is an object of the present invention to provide a paper web having excellent formation and with a capacity of forming particularly strong bonds.

[0009] The present invention is based on the idea of forming the base paper from a mixture of mechanical and chemical pulp, the chemical pulp used comprising a chemical softwood pulp incorporating in combination a large ScottBond strength and an elastic modulus which is relatively small for chemical softwood pulp. Preferably the elastic modulus is less than 6000 N/mm², when the ScottBond-strength of the chemical pulp is 400 J/m². Thus, a paper produced from a mixture of mechanical pulp and chemical pulp will simultaneously have high ScottBond strength and large toughness.

[0010] More specifically, the solution according to the present invention is mainly characterized by what is stated in the characterizing part of claim 1.

[0011] Considerable advantages are obtained by the present invention. Thus, the pulp produced according to the invention has at the same amount of surface bonding, i.e. at the same light scattering, a better bonding strength than comparative pulps. The present base paper can therefore be used for production of double-coated fine papers which in particular require great bonding strength of the base paper. Other fiber components whose internal bond strength in itself is not sufficient can be incorporated into the base paper. As a specific example reference can be made to the manufacture of fine paper from mixtures of aspen groundwood and chemical softwood pulp, whereby a strong paper is obtained as a finished product, said paper having good brightness and opacity and a very smooth surface. Thanks to the good bonding strength of the chemical softwood pulp, aspen groundwood can be used even in amounts up to 20 to 60 % of the dry matter of the pulp.

[0012] The technical solution according to the present invention comprises using a chemical pulp which has been produced by chemical pulping which will protect the fibers, whereby their strength remains good. The cooking should

be selective in the sense that it selectively removes lignin and spares the carbohydrates of the fiber. In connection with the present invention it has been found that these objects can be obtained by using batch cooking, a particularly preferred embodiment comprising extended batch cooking (Superbatch cooking).

5 [0013] As regards the strength of the chemical pulp, the pulping method is not as such a sufficient criterion, but the chemical pulp produced according to the invention should have enough bonds between the fibers. In connection with the present invention it has been found that by bleaching softwood pulp produced by batch cooking with TCF bleaching comprising bleaching stages with peroxide and ozone particularly good strength properties are obtained. Said oxidizing chemicals form carboxylic groups on the fibers and these groups improve the strength of the bleached pulp.

10 [0014] The importance of the acid groups for forming bonds between the fibers has been discussed in Barzyk, D. et al. Journal of Pulp and Paper Science, 23 (1997) J59-J61. According to that article the bonding strength is based on carboxylic groups. In the present invention it has, however, been found that it is not only the amount of acid groups that is decisive, but the conditions of the cook and the bleaching sequences are also of importance.

[0015] As discussed above, when attempts are made to regulate the properties of the pulp by beating, i.e. when the ScottBond is raised by a high degree of beating, the chemical pulp and, e.g., hardwood groundwood, will get very different elastic moduli (chemical pulp gets very high stiffness), which is undesirable as far as the toughness of a mixture produced of these pulps is concerned. This problem is not encountered in the present invention. For this reason, by means of the present invention, a mixture of hardwood groundwood and chemical pulp is obtained which is excellent as a base paper of fine papers.

20 [0016] According to a preferred embodiment, the chemical pulp used for preparing a base paper is produced by a cooking method known as a modified batch-type cook (Superbatch Cook). This cook is described in the literature [cf. for example Malinen, R. Paperi ja Puu (Paper and Timber), 75 (1993) 14-18]. The cook in question is a modified cooking method which utilizes an alkaline cooking liquor just as the sulphate cook, but wherein delignification has been enhanced so that the kappa number of the chemical pulp is lowered without a significant reeduction of viscosity. Typically with a Superbatch process, pulp is cooked to a kappa number of 20 or less.

25 [0017] According to a preferred embodiment of the present invention, a softwood pulp produced by batch cooking is bleached with TCF bleaching. The following examples of suitable bleaching sequences can be mentioned:

(Q)-O-Z-P-Z-P

30

(Q)-O-Z-E-P_n

35

O-(Q)-Z-E-P-Z-E-P

O-Z-(Q)-P_n

40

O-X-Z-P_n

O = oxygen treatment

P = peroxide treatment

45 P_n = several successive peroxide treatment stages

E = alkali step

Q = treatment with complexing agent

X = enzyme treatment

50 [0018] An acid pretreatment at elevated temperature (an A stage) can be performed between the oxygen delignification (O-stage) and a bleaching step carried out with an oxidizing chemical (i.e. a Z-stage).

[0019] It is particularly preferred to carry out the bleaching of the pulp with two ozone stages and at least two peroxide stages. Between the stages carried out with oxidizing chemicals, it is possible to extract the pulp during various alkaline stages (such as E and E0) and/or to wash it with water.

55 [0020] Following the above-described treatment a pulp is obtained having an internal bond strength which is better than that of comparative pulps. It typically contains at least 40 mmol carboxylic acid groups/kg dry pulp. Preferably the elastic modulus of the chemical pulp used according to the present invention is below 6000 N/mm², in particular below 5000 N/mm² when ScottBond strength is 400 J/m².

[0021] As mentioned above, the base paper can be produced from chemical pulp by combining it with aspen ground-wood, by slushing the obtained fibrous base material, by forming a web from the stock and by drying the web on a paper machine in order to form a base paper. Generally, the pulp can be produced from any mechanical pulp made of a tree of the *Populus* family. Suitable species are, for example, *P. tremula*, *P. tremuloides*, *P. balsamea*, *P. balsamifera*, *P. trichocarpa* and *P. heterophylla*. A preferred embodiment comprises using aspen (trembling aspen, *P. tremula*; an aspen known as Canadian aspen, *P. tremuloides*), or aspen varieties known as hybride aspens produced from different base aspens by hybridizing as well as other species produced by recombinant technology, or poplar. It is preferred to use groundwood (GW), pressure groundwood (PGW) or thermomechanical pulp (TMP) manufactured from aspen, hybride aspen or poplar.

[0022] Preferably the mechanical aspen pulp contains about 10 to 20 % of +20... +48 mesh fibers, which confer mechanical strength to the pulp. In order to maximize light scattering, the portion of + 100, +200 and -200 fractions should be as large as possible. Preferably they stand for distinctly more than 50 % of the whole pulp. In particular their proportion of the whole pulp is over 70 %, preferably over 80 %. On the other hand, the amount of the smallest fraction, i.e. the -200 mesh, should not be too large, because then dewatering on the paper machine would become more difficult. Preferably the proportion of this fraction is smaller than 50 %, in particular 45 % or less.

[0023] Due to the excellent mechanical properties of the pulp according to the present invention the proportion of the mechanical pulp can be even up to 70 weight-% of the dry matter of the stock without the strength of the paper essentially suffering. Typically, the proportion of the mechanical pulp is at least 20 % and in particular it is 30 to 60 weight-%.

[0024] Based on what is stated above, according to the invention the composition of a particularly preferred base paper is the following: 30 to 60 weight-% of the fibrous matter comprises mechanical pulp produced from aspen and 70 to 40 weight-% comprises softwood chemical pulp. The ScottBond strength of the chemical softwood (in particular pine) pulp is at least 400 J/m² at a light scattering coefficient of 22 m²/kg and it contains at least 40 mmol carboxylic acid groups/kg dry pulp.

[0025] From the base paper according to the present invention it is possible to produce high-quality fine paper by coating it preferably twice, the first coating for example being carried out by a method known as the film press method, and the second coating is performed by blade coating. The amount of coating colour applied to the web by the film press method is typically about 5 to 50 g coating colour/m², whereas the corresponding amount for doctor blade coating is 10 to 60 g coating colour/m². The indicated amounts of coating have been calculated from the dry matter of the coating colour.

[0026] Next, the invention will be examined more closely with the aid of a detailed description and with reference to the attached drawings and working examples.

Figure 1 compares the pulps disclosed in the examples; the ScottBond strength is indicated on the y axis as a function of the light scattering coefficient,

Figure 2 indicates the ScottBond strengths of three mixed sheets as a function of the light scattering coefficient, and

Figure 3 contains a comparison of the elastic moduli of four chemical pulps as a function of internal bond strength.

[0027] The following measurement standards have been used in the examples:

- ISO brightness of the chemical pulp: SCAN-C'M 11 and SCAN-P3
- light scattering coefficient: SCAN-C 27
- ScottBond strength: Tappi T833
- brightness: SCAN-P3:93 (D65/10°)
- opacity: SCAN-P8:93 (C/2)
- surface coarseness: SCAN-P76:95
- Bendtsen coarseness: SCAN-P21:67
- gloss: Tappi T480 (75°) and T653 (20°)
- elastic modulus measurement: SCAN-P 38 (strip size and tensile velocity)

[0028] For measurement of the elastic modulus the sheet was prepared and the drying was carried out according to standard SCAN-C 26.

Example 1

Internal bond strength of chemical pulps

[0029] The ScottBond strength of sheet produced from softwood chemical pulp is influenced by the extent of bonding

surface between the fibres and the strength of the bonds. The amount of the bonding surface is, on its part, strongly dependent on the degree of beating of the chemical pulp used in sheetmaking. When beating is increased the bonding area and at the same time the bonding strength are increased. To make it possible to compare bond strengths, in this example the internal bond strengths of different chemical pulps are compared by examining them as a function of the light scattering coefficient in the same way as in the article by Barzyk et al. Journal of Pulp and Paper Science, 23 (1997) J59-J61, Figures 3 and 4, already referred to above. It is conceivable that with chemical softwood pulps the light scattering coefficient is a measure of the amount of bonding surface of the fibers, the greater the amount of bonding surface the smaller the light scattering coefficient.

[0030] In this test the internal bond strength and light scattering coefficient of chemical pulps have been modified by beating the pulps in an Escher-Wyss-refiner at various energy amounts of 0 to 200 kWh/ton. The specific edge load of the beating was 3 Ws/m. The results are indicated in Figure 1. In that Figure, the curve extending to a higher level at the same amount of bonding surface, i.e. light scattering, stands for an increased bonding strength.

[0031] Graphs 1 to 3 depict cellulosic pulps produced by a continued batch cooking (SuperBatch) which have been subjected to chlorine-free bleaching (TCF) by using two ozone and two peroxide stages (ZPZP). Graphs 4 and 5 depict a pulp produced by a continuous cooking method, which also has been subjected to chlorine-free bleaching (TCF) by using one ozone and one peroxide stage (ZP). The cooking result is, compared to the above mentioned batch cooking, more heterogenous and weaker fibers are produced. The fiber collapses more easily, and it loses its light scattering coefficient which moves the curve to the left. The pulps produced by both methods 1 to 3 and 4 and 5 contain at least approximately an equal amounts of carboxylic acid groups (41 - 47 mekv/kg and 42 - 46 mekv/kg, respectively).

[0032] Graphs 6 to 9 show pulps which have been subjected to a bleaching without elemental chlorine (ECF bleaching). The starting material of cooking 6 was a raw material obtainable in the north of Finland. It comprises small size fibers which give a large specific surface (m²/g fiber) and, therefore, its light scattering coefficient is good. The concentration of carboxylic acid groups was 34 mekv/kg. The raw material of cooking 7 was obtained from Eastern Finland and the chemical pulp had been produced by batch cooking. Graphs 8 and 9 represent the internal bonding strength of pulps produced by continuous cooking and bleached by ECF bleaching. The concentration of carboxylic groups was 27 to 34 mekv/kg. The graphs show that pulps 1 to 3 give greater values for the bonding strength than the other pulps at the same light scattering coefficient. The differences become more pronounced when the pulp have been subjected to extended beating.

[0033] Next, three of the afore-mentioned pulps were selected for a sheet forming test. Although the pulps were not from the same batches as above, pulp A corresponded to pulps 1 to 3, pulp B corresponded to pulp 6 and pulp C corresponded to pulp 7. The pulps were refined in a laboratory Valley beater so that the degree of beating (drainage) was CSF 380 ml. Then sheets were produced from the pulps so that in each test point the sheets contained 60 % chemical pulp and 40 % aspen PGW pulp (aspen of *Populus* family).

[0034] When the bonding strengths of the mixed sheets vs. light scattering coefficient now were examined, a result according to Figure 2 was obtained. Even if the differences are rather small, it is apparent that chemical pulp A gives a better result than pulps B and C. The trend is the same as for pure pulp sheets; in other words: by the combination of batch cooking and TCF bleaching according to the present invention a better bonding strength is obtained than for the comparative pulps, even if these separately include the partial elements of the invention.

[0035] Finally, an analysis was made to determine how the elastic modulus develops as a function of ScottBond strength. This test included pulps from three production batches (A1, A2 and A3), which corresponded to pulps 1 to 3 of Figure 1, and a pulp sample D which corresponded to pulps 8 and 9 of Figure 1. Pulp samples A1 and A2 had been refined to different beating degrees in a Escher-Wyss refiner and samples A3 and D again in a Valley beater. Figure 3 shows that the elastic modulus of pulp A was smaller than for pulp D, if the comparison is carried out at the same ScottBond strength. Thus, it can be expected from the pulp A according to the invention that it gives a smaller elastic modulus than D and, accordingly, that a paper produced from pulp A is less brittle. In other words, the paper is tougher than a paper made from pulp D. The superiority of pulp A is pronounced when the pulps are beaten to a high degree of beating in order to obtain good ScottBond strength.

Example 2

Production of a fine paper containing aspen PGW

[0036] A base paper was produced from a mechanical aspen pulp (GW) and chemical pine pulp, which were mixed at a weight ratio of 40 to 60. Ground calcium carbonate was added as a filler to the suspension in an amount of about 10 % of the fibrous material.

[0037] The base paper was produced on a gap former. The properties of the base paper were the following:

grammage	53.3 g/m ²
bulk	1.45 cm ³ /g
opacity	88 %
brightness	82.5 %
coarseness	240 ml/min
porosity	170 ml/min
filler content	12 %

[0038] Comparative test carried out in connection with the invention have shown that the grammage of the base paper is at least 10 % smaller than that of a base paper produced entirely from a bleached chemical pulp and having the corresponding opacity and brightness.

[0039] In order to produce the fine paper from the above-described base paper it was coated twice, first with the film press method and then with doctor blade coating.

[0040] A calcium carbonate pigment having the particle size distribution shown in Table 1 was used in the coating colours:

Table 1.

Particle size distribution of the carbonate pigment	
Max. particle size [µm]	Cumulative proportion of weight
5	99
2	95
1	70
0.5	35
0.2	10

[0041] The coating colour was produced in a manner known *per se* by mixing together the pigment, the binder and the other additives. The dry matter content of the precoat colour was 60 % and of the surface coating colour 61 %. The above described colours were used for coating the afore-mentioned base paper in the following conditions:

[0042] Precoat by the film press method: 9 g/m² per side; and the surface coating at a doctor blade station: 10.5 g/m² per side at a speed of 1500 m/min. The coated paper was super-calendered.

[0043] The properties of the end products were determined and compared to those of two commercially available finer papers, viz. Lumiart (Enso) and Nopacoat (Nordland Papier). The results will appear from Table 2:

Table 2.

Optical properties of a double-coated fine paper			
	Paper according to the invention	Lumiart	Nopacoat
Grammage [g/m ²]	80	100	99
Bulk	0.85	0.83	0.78
Opacity [%]	94	92.7	92.6
Brightness [%]	94	91	96.7
Smoothness pps 10 [µm]	0.8	1.2	0.8
Gloss [%]	73	66	71

[0044] Table 2 shows that the properties of a fine paper produced by the invention are better in all respects than those of comparative papers having corresponding bulk and grammage. Thus the yield gain on equal level of opacity is over 20 %.

[0045] The ScottBond bonding strength of the fine paper prepared according to the Example was 306 J/m². This is also fully comparable to the strength of a traditional fine paper comprising only chemical pulp. Even if the internal bonding strength of aspen PGW is inferior to that of e.g. chemical birch pulp, the present invention has provided a

paper which is strong enough for use as a fine paper.

Claims

5

10

15

20

25

30

35

40

45

50

55

1. A process for producing base paper for coated fine papere, **characterized** in that the base paper is produced from a mixture of mechanical pulp and chemical pulp, whereby the chemical pulp used comprises softwood pulp having a elastic modulus which is close to the elastic modulus of elasticity of the mechanical pulp and exhibiting a large bonding strength.
2. The process according to claim 1, **characterized** by using a chemical softwood pulp, which can be used for producing a sheet having a elastic modulus of less than 6000 N/mm² when the bonding strength is 400 J/m².
3. The process according to claim 1 or claim 2, **characterized** by using mechanical pulp produced from hardwood and bleached softwood cellulosic pulp produced by batch cooking.
4. The process according to claim 2 or claim 3, **characterized** by using softwood cellulosic pulp bleached with ozone and peroxide.
5. The process according to claim 4, **characterized** by using softwood cellulosic pulp bleached with at least two ozone stages and at least two peroxide stages.
6. The process according to any of the preceding claims, **characterized** by using a cellulosic pulp produced by the SuperBatch process.
7. The process according to any of claims 1 to 6, **characterized** by using cellulosic pulp having a ScottBond strength amounting to at least 400 J/m² at a light scattering coefficient of 22 m²/kg and containing over 40 mequivalents of carboxylic acid groups per kg of dry pulp.
8. The process according to any of the preceding claims, **characterized** by using a cellulosic pulp having a brightness of more than 82, preferably over 85 and in particular over 88.
9. The process according to any of the preceding claims, according to which process
 - a stock is formed from a fibrous raw material,
 - the stock is formed into a web, and
 - the web is dried in order to form a base paper,**characterized by**
 - forming the stock from mechanical pulp produced from wood material from the *Populus* family and from TCF bleached softwood cellulosic pulp produced by batch cooking, the amount of mechanical pulp being 20 to 70 weight- % and that of the bleached softwood cellulosic pulp being 80 to 30 weight- % of the dry matter of the suspension.
10. The process according to claim 9, **characterized** in that 30 to 60 weight- % of the dry matter of the stock is formed by mechanical pulp and 70 to 40 weight- % by softwood cellulosic pulp.
11. The process according to claim 9 or 10, **characterized** in that the mechanical pulp is produced from *P. tremula*, *P. tremuloides*, *P. balsamea*, *P. balsamifera*, *P. trichocarpa* or *P. heterophylla*.
12. The process according to claim 11, **characterized** in that the mechanical pulp is pressure ground wood.
13. Base paper for fine papers, **characterized** in that 30 to 60 weight-% of its fibrous material consists of a mechanical pulp produced from aspen and 70 to 40 weight-% consists of chemical softwood pulp, the latter having a ScottBond strength amounting to at least 400 J/m² at a light scattering coefficient of 22 m²/kg and containing over 40 mequivalents of carboxylic acid groups per kg of dry pulp.

EP 0 908 558 A1

14. A process for producing a coated fine paper, **characterized** in that a base paper produced according to any of claims 1 to 11 or a base paper according to claim 12 is coated with two coating colour layer, the first coating being conducted by the film press method and the second by doctor blade coating.

5 15. The process according to claim 14, **characterized** by applying on the web 5 to 50 g coating colour/m² by the film press method and 10 to 60 g coating colour/m² by doctor blade coating, the coating weights being calculated based on the dry matter of the coating colour.

10

15

20

25

30

35

40

45

50

55

Chemical pulp sheets

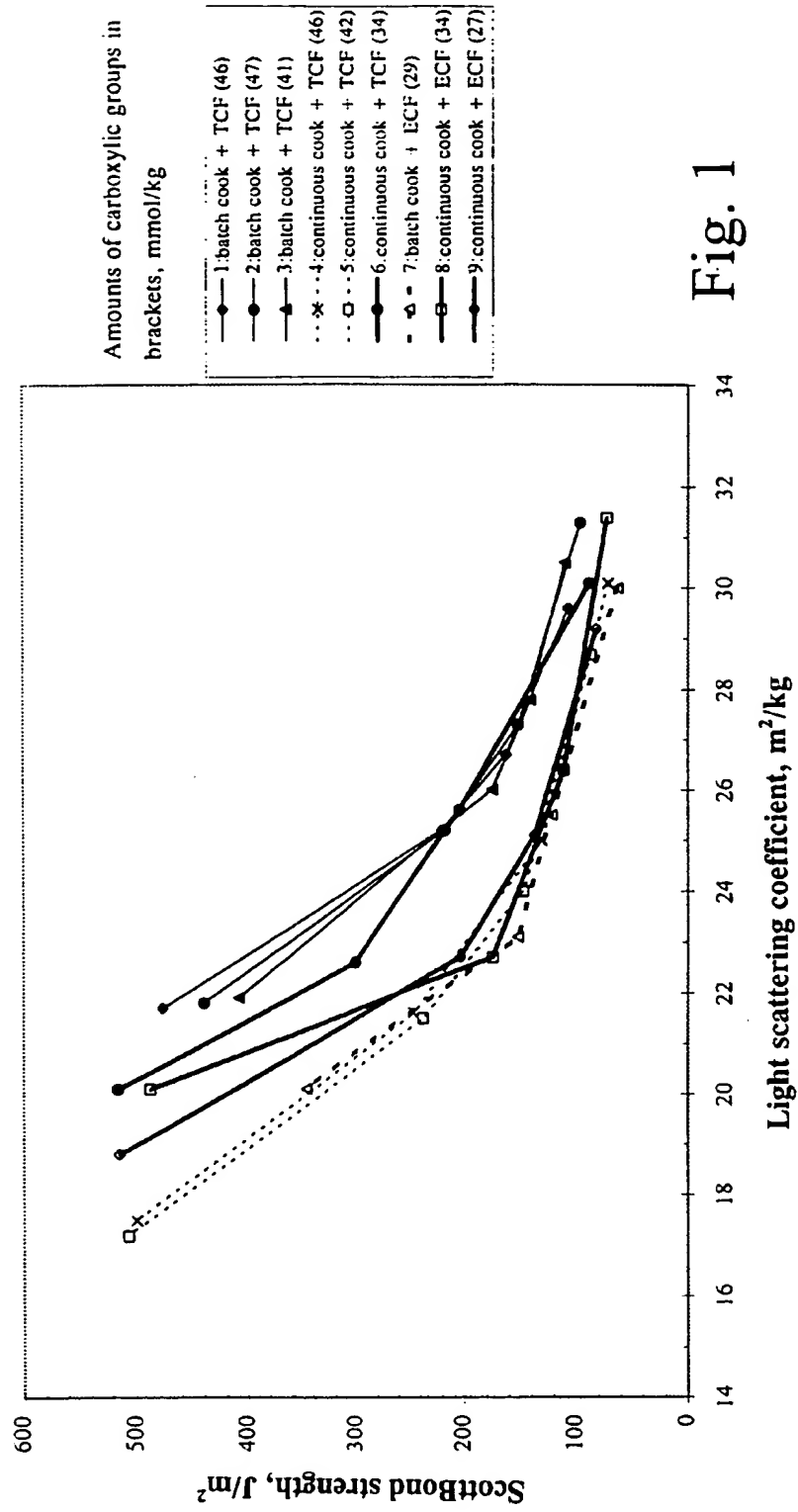


Fig. 1

Sheets

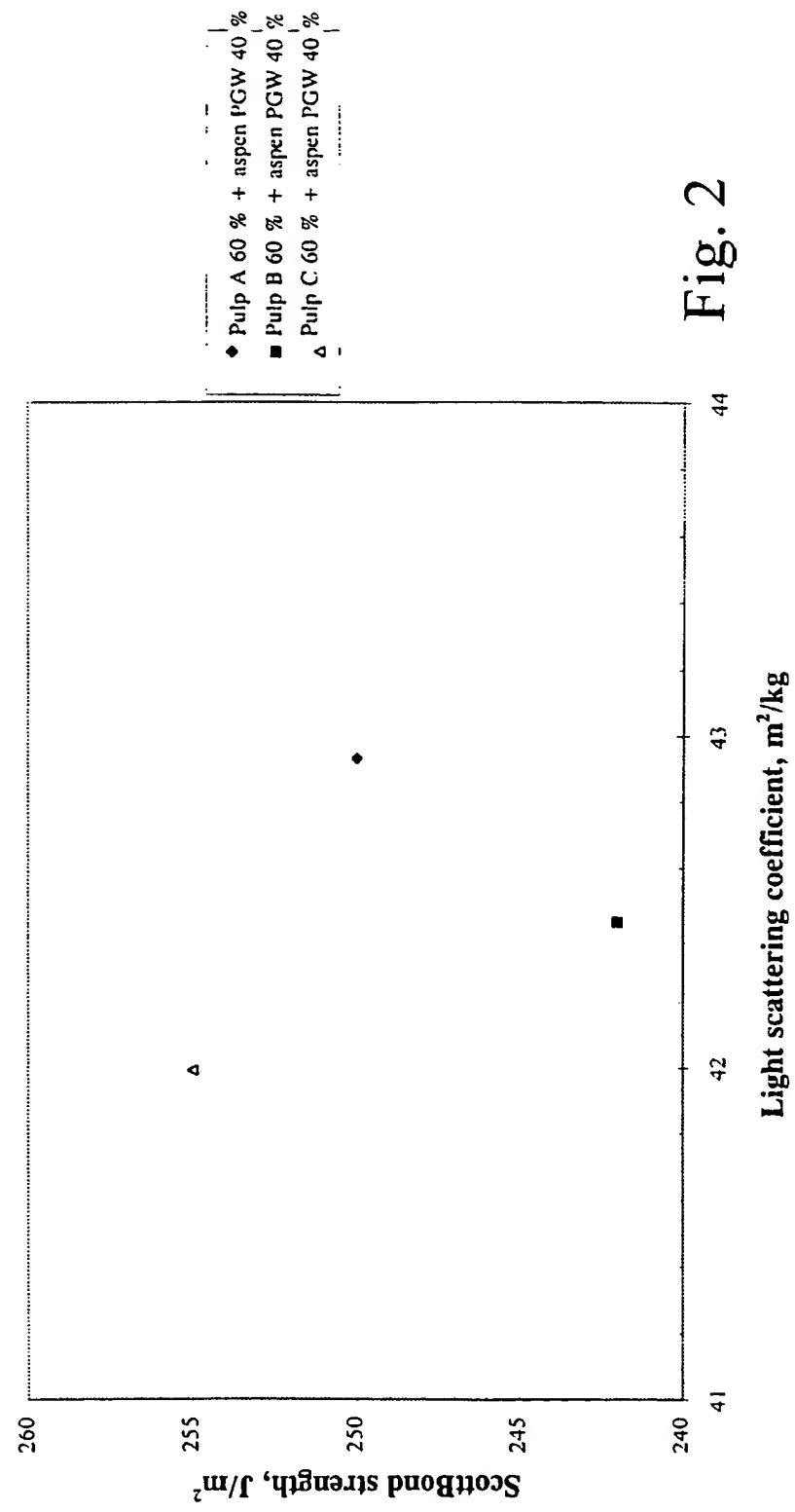


Fig. 2

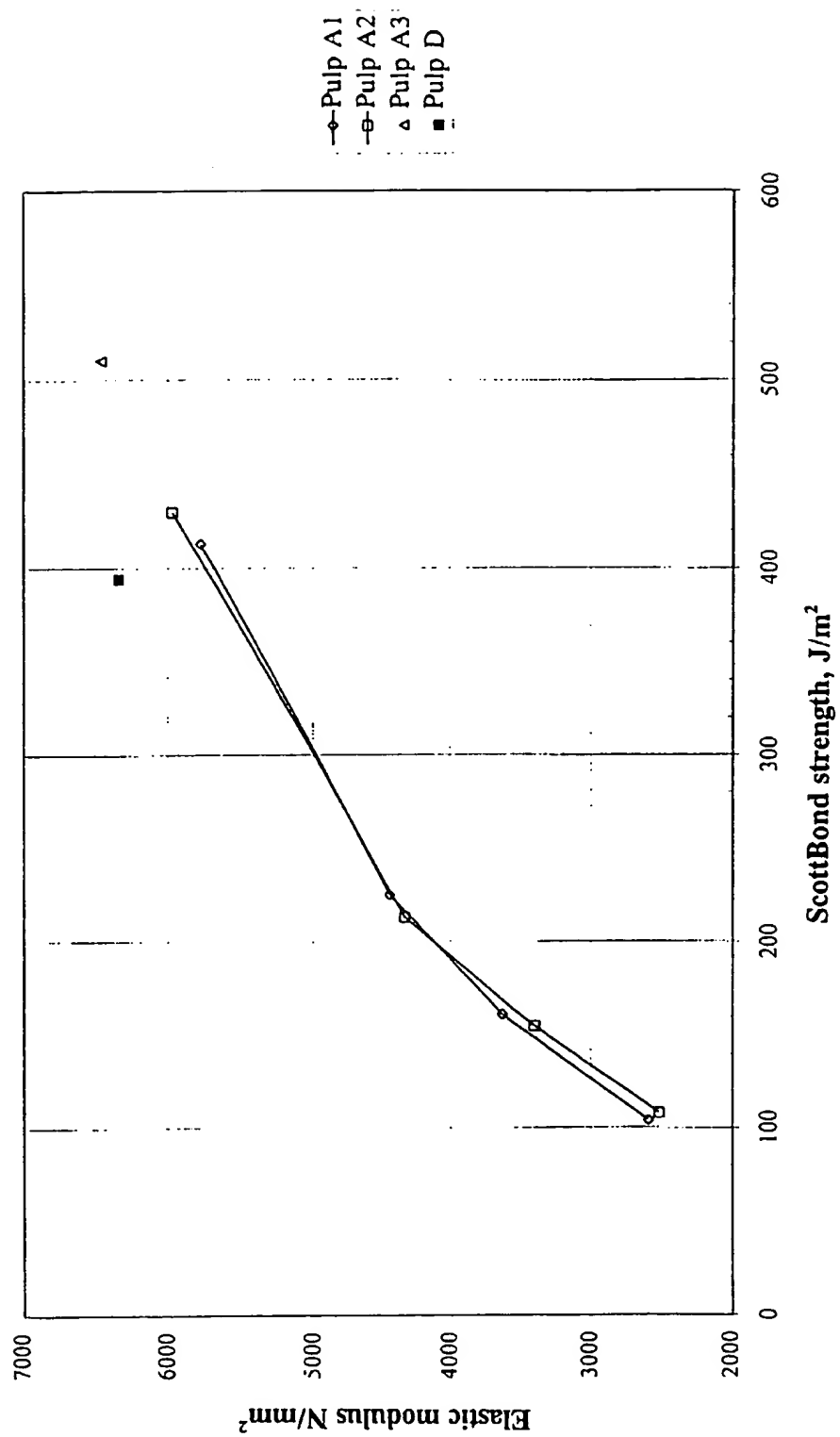


Fig. 3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 98 66 0095

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
Y	DATABASE PAPERCHEM THE INSTITUTE OF PAPER SCIENCE AND TECHNOLOGY, ATLANTA, GA, US HOYDAHL, H.-E: "THERMOMECHANICAL PULP - FASHIONABLE OR REALISTIC FIBER ALTERNATIVE" XP002086903 * abstract *	1,9-11	D21H11/10 D21H23/72
A	& NORSK SKOGIND. 29, NO. 12: 323-328 (DEC. 1975). 'NORW.;ENGL. SUM.!, ---	12,13	
Y	DATABASE PAPERCHEM THE INSTITUTE OF PAPER SCIENCE AND TECHNOLOGY, ATLANTA, GA, US KARENLAMPI, P.: "Tensile Strength of a Mixture of Two Pulps" XP002087047 * abstract * & J. PULP PAP. SCI. 21, NO. 12: J432-436 (DECEMBER 1995). 'ENGL.;FR. SUM.!, ---	1	
Y	DATABASE PAPERCHEM THE INSTITUTE OF PAPER SCIENCE AND TECHNOLOGY, ATLANTA, GA, US PISAREVSKAYA, E. A. ET AL: "PRODUCTION OF BLEACHED ASPEN GROUNDWOOD AT THE ZHIDACHEV MILL" XP002086900 * abstract * & BUMAZH. PROM. NO. 1: 17-18 (JAN. 1981). 'RUSS.!, ---	1,9-11	TECHNICAL FIELDS SEARCHED (Int.Cl.6) D21H
Y	US 4 298 652 A (SUZUKI KUZUHIKO ET AL) 3 November 1981 * the whole document *	1	
A	---	12-15	
	---	-/-	
The present search report has been drawn up for all claims			
Place of search MUNICH		Date of completion of the search 8 December 1998	Examiner Nestby, K
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X: particularly relevant if taken alone Y: particularly relevant if combined with another document of the same category A: technological background O: non-written disclosure P: intermediate document</p> <p>T: theory or principle underlying the invention E: earlier patent document, but published on, or after the filing date D: document cited in the application L: document cited for other reasons &: member of the same patent family, corresponding document</p>			

EPO FORM 1503 03.92 (P04/C01)



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 98 66 0095

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
A	GB 2 189 714 A (YHTYNEET PAPERITEHTAAT OY) 4 November 1987 * page 1, line 24 - line 39 *	14,15	
A	DE 44 11 987 A (FELDMUEHLE AG STORA) 12 October 1995 * page 2, line 44 - page 3, line 4 *	14,15	
A	US 5 340 611 A (KUSTERMANN MARTIN ET AL) 23 August 1994 * the whole document *	14,15	
			TECHNICAL FIELDS SEARCHED (Int.Cl.6)
The present search report has been drawn up for all claims			
Place of search MUNICH		Date of completion of the search 8 December 1998	Examiner Nestby, K
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

EPO FORM 1600 (03/96) (P0401)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 98 66 0095

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report.
The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

08-12-1998

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4298652 A	03-11-1981	JP 55152895 A	28-11-1980
		JP 63027480 B	03-06-1988
		AU 533973 B	22-12-1983
		AU 5831680 A	13-11-1980
		CA 1147619 A	07-06-1983
		DE 3017763 A	20-11-1980
		FR 2456162 A	05-12-1980
		GB 2050868 A,B	14-01-1981
GB 2189714 A	04-11-1987	FI 861807 A	31-10-1987
		DE 3714498 A	05-11-1987
		FR 2598160 A	06-11-1987
		FR 2637623 A	13-04-1990
		GB 2205051 A,B	30-11-1988
		JP 62263397 A	16-11-1987
		NL 8701022 A	16-11-1987
		SE 504900 C	26-05-1997
DE 4411987 A	12-10-1995	SE 8701747 A	31-10-1987
		AT 171743 T	15-10-1998
		CA 2187182 A	19-10-1995
		DE 59503785 D	05-11-1998
		WO 9527824 A	19-10-1995
		EP 0754260 A	22-01-1997
		FI 963942 A	02-10-1996
		JP 9511552 T	18-11-1997
US 5340611 A	23-08-1994	DE 4002256 A	01-08-1991
		CA 2035061 A	27-07-1991
		DE 59003011 D	11-11-1993
		EP 0438743 A	31-07-1991
		ES 2044390 T	01-01-1994
		FI 905757 A	27-07-1991
		JP 4313360 A	05-11-1992
		NO 176676 B	30-01-1995
		US 5179909 A	19-01-1993

EPO FORM P449

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82